

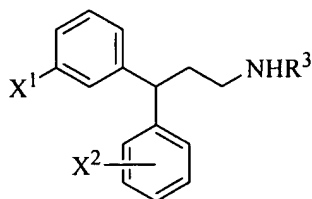
**IN THE CLAIMS:**

Claims 5 and 25-27 are amended herein. New claims 28 and 29 have been added. All claims currently pending and under consideration in the referenced application are shown below.

**Listing of Claims:**

1-4. (Canceled)

5. (Currently amended) A method of treating a patient for depression comprising administering to ~~said~~ the patient an effective amount of a compound having the following chemical structure:



wherein X<sup>1</sup> is either -Br, -Cl, -F, -I, -CF<sub>3</sub>, alkyl, -OH, -OCF<sub>3</sub>, -O-alkyl, or -O-acyl;

X<sup>2</sup> is either -Br, -Cl, -F, -I, -CF<sub>3</sub>, alkyl, -OH, -OCF<sub>3</sub>, -O-alkyl, or -O-acyl; ~~and~~

R<sup>3</sup> is either -H or -CH<sub>3</sub>; and

~~or a~~ pharmaceutically acceptable salts thereof.

6. (Original) The method of claim 5, wherein X<sup>1</sup> is -F, -Cl, -OCF<sub>3</sub> or -CF<sub>3</sub> and X<sup>2</sup> is either 2-OCH<sub>3</sub>, 2-CH<sub>3</sub>, 3-F, 3-CF<sub>3</sub>, or 4-CF<sub>3</sub>.

7-20. (Canceled)

21. (Previously presented) The method of claim 5, wherein X<sup>1</sup> and X<sup>2</sup> are F, and R<sup>3</sup> is -H.

22. (Previously presented) The method of claim 21, wherein X<sup>2</sup> is at the 3-position.

23. (Previously presented) The method of claim 5, wherein  $X^1$  and  $X^2$  are F, and  $R^3$  is  $-CH_3$ .

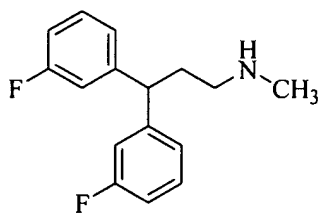
24. (Previously presented) The method of claim 23, wherein  $X^2$  is at the 3-position.

25. (Currently amended) The method of claim 5, wherein ~~said the~~ compound is active at a serotonin reuptake site and at a N-methyl-D-aspartate (NMDA) receptor.

26. (Currently amended) The method of claim 5, wherein ~~said the~~ compound has an NMDA receptor  $IC_{50}$  of about 50 nM to about 1  $\mu$ M.

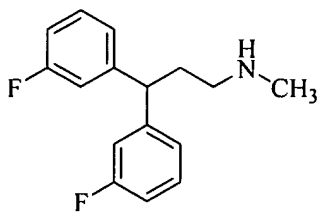
27. (Currently amended) The method of claim 26, wherein ~~said the~~ compound has an NMDA receptor  $IC_{50}$  of about 100 nM to about 800 nM.

28. (New) The method of claim 5, wherein the compound has the chemical structure:



and pharmaceutically acceptable salts thereof.

29. (New) The method of claim 27, wherein the compound has the chemical structure:



and pharmaceutically acceptable salts thereof.